

Rec'd PCT/PTO 13 JUL 2005

10/542187

(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property  
Organization  
International Bureau



(43) International Publication Date  
5 August 2004 (05.08.2004)

PCT

(10) International Publication Number  
WO 2004/065920 A2

(51) International Patent Classification<sup>7</sup>: G01J  
(21) International Application Number:  
PCT/US2004/001144  
(22) International Filing Date: 16 January 2004 (16.01.2004)  
(25) Filing Language: English  
(26) Publication Language: English  
(30) Priority Data:  
60/440,887 17 January 2003 (17.01.2003) US

(81) Designated States (*unless otherwise indicated, for every kind of national protection available*): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW.

(84) Designated States (*unless otherwise indicated, for every kind of regional protection available*): ARIPO (BW, GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European (AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).

(71) Applicant (*for all designated States except US*): GRIF-FIN ANALYTICAL TECHNOLOGIES, INC. [US/US]; 3000 Kent Avenue, West Lafayette, IN 47906 (US).

(72) Inventors; and

(75) Inventors/Applicants (*for US only*): GROSSEN-BACHER, John, W. [US/US]; 1309 Ridgeway Ave., Lafayette, IN 47904 (US). PATTERSON, Garth, E. [US/US]; 802 S. Brackney, Brookston, IN 47923 (US).

(74) Agent: HYTA, Robert, C.; Wells St. John, P.S., 601 W. 1st Avenue, Suite 1300, Spokane, WA 99201 (US).

Published:

— without international search report and to be republished upon receipt of that report

*For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.*

(54) Title: MASS SPECTROMETER ASSEMBLIES MASS SPECTROMETRY VACUUM CHAMBER LID ASSEMBLIES AND MASS SPECTROMETER OPERATIONAL METHODS

(57) Abstract: One embodiment of a mass spectrometer assembly includes a base configured to define at least a portion of a vacuum chamber volume; a mass separator component within the vacuum chamber volume; a lid coupled to the mass separator component and configured to be removably operably coupled with respect to the base; and wherein the lid is configured to be positioned in a first operable position to form a hermetical seal with the base and provide the mass separator component within the vacuum chamber volume and a second operable position wherein at least a portion of the lid is spaced from the base and the mass separator component is at least partially removed from within the vacuum chamber volume. Mass spectrometer operational methods are provided that can include at least partially removing the lid from the base, wherein the removing of the lid also at least partially removes the mass separator component from the vacuum chamber volume; and inspecting the mass separator component with the mass separator component removed from the vacuum chamber volume.

WO 2004/065920 A2

**DESCRIPTION****MASS SPECTROMETER ASSEMBLIES,  
MASS SPECTROMETRY VACUUM CHAMBER LID ASSEMBLIES, AND MASS  
SPECTROMETER OPERATIONAL METHODS****CLAIM FOR PRIORITY**

This application claims priority to United States provisional patent application Serial Number 60/440,887 filed January 17, 2003, entitled "Interchangeable Mass Spectrometer Inlet/Ionization Source", the entirety of which is hereby incorporated by reference.

**TECHNICAL FIELD**

The present disclosure relates generally to chemical analysis and more particularly to mass spectrometer assemblies, mass spectrometry vacuum chamber lid assemblies, and mass spectrometer operational methods.

**BACKGROUND ART**

Characterization of compounds utilizing mass spectrometry and varying sources of ionization is well accepted in the field of analytical chemistry as a technique that allows for the further elucidation of analytes and their specific chemistries. However, mass spectrometer instrumentation is costly and because most labs are unable to configure many instruments with unique ionization sources, analysts are typically required to configure one instrument with a single source and then reconfigure with different sources as analysis dictates. This change-out between sources can be problematic, particularly since mass spectrometer instrumentation must be configured under a vacuum and nanogram quantities of contaminant materials can provide background noise rendering the instrument practically useless.

**SUMMARY**

Mass spectrometer assemblies are provided that can include in one embodiment: a base configured to define at least a portion of a vacuum chamber volume within which at least some operations may be performed with respect to mass spectrometry; a mass separator component configured to perform at least some operations with respect to mass spectrometry within the vacuum chamber volume; a lid coupled to the mass separator component and configured to be removably operably coupled with respect to the base; and wherein the lid is configured to be positioned in a first operable position to

form a hermetical seal with the base and provide the mass separator component with the vacuum chamber volume and a second operable position wherein at least a portion of the lid is spaced from the base and the mass separator component is at least partially removed from the vacuum chamber volume.

5        Mass spectrometry vacuum chamber lid assemblies are provided that can include, in one embodiment, a body having an interior surface coupled to a mass separator component, wherein the body is configured to at least partially define a volume partially surrounding the mass separator component when the body is hermetically sealed to a housing of a vacuum chamber assembly, wherein the body is further  
10        configured to be removable from the vacuum chamber volume to at least partially remove the mass separator component from the vacuum chamber volume.

      Mass spectrometer operational methods are provided that can include, in one embodiment: providing a mass spectrometry assembly comprising a base and a lid, the base and lid substantially defining a vacuum chamber volume when the lid is affixed to  
15        the base in a position operable to perform at least some operations with respect to mass spectrometry, wherein a mass separator component is coupled to the lid and occupies a portion of the vacuum chamber volume in the position; first performing mass analysis using the mass spectrometry assembly in the position; after the first performing, at least partially removing the lid from the base, wherein the at least partially removing of the lid  
20        also at least partially removes the mass separator component from the vacuum chamber volume; inspecting the mass separator component with the mass separator component removed from the vacuum chamber volume; returning the lid to the position after the inspecting; and second performing mass analysis using the mass separator after the returning.

25

#### **BRIEF DESCRIPTION OF THE DRAWINGS**

      Preferred embodiments of the invention are described below with reference to the following accompanying drawings.

30        Fig. 1 is an illustrative representation of a mass spectrometry assembly according to an embodiment.

      Fig. 2 is a block diagram of mass spectrometry components according to an embodiment.

      Fig. 3a is an illustrative representation of a mass spectrometry assembly according to an embodiment.

35        Fig. 3b is an illustrative representation of a mass spectrometry assembly according to an embodiment.

Fig. 3c is an illustrative representation of a mass spectrometry assembly according to an embodiment.

Fig. 3d is an illustrative representation of a mass spectrometry assembly according to an embodiment.

5 Fig. 4 is an isometric view of a mass spectrometry assembly according to an embodiment.

Fig. 5a is a plan view of a mass spectrometry assembly according to an embodiment.

10 Fig. 5b is a plan view of the mass spectrometry assembly of Fig. 5a according to an embodiment.

Fig. 5c is a side view of the mass spectrometry assembly Figs. 5a and 5b according to an embodiment.

Fig. 6 is a plan view of a mass spectrometry assembly according to an embodiment.

15 Fig. 7 is an illustrative representation of a mass spectrometry assembly according to an embodiment.

#### **DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS**

20 At least some embodiments provide mass spectrometry assemblies and mass spectrometer operational methods. Exemplary configurations of these assemblies and methods are described with reference to figures 1-7.

Referring first to Fig. 1, a mass spectrometry assembly 10 is shown that comprises a base 12 and a lid 14. Together lid 14 and base 12 can define a volume 16. In an exemplary embodiment lid 14 and base 12 define a vacuum chamber housing of mass spectrometry assembly 10. The vacuum chamber housing can define volume 16 which can be referred to as a vacuum chamber volume in embodiments wherein a vacuum is provided. A vacuum source (not shown) can be in fluid connection to base 12 and provide vacuum to volume 16.

30 Base 12 can be constructed of a single structure or can be constructed of multiple components. Exemplary components include walls 17 and a bottom 18. In the exemplary configuration of Fig. 1, walls 17 may be a continuous wall of a cylinder. Other geometries or arrangements of base 12 are possible. The structure and or components of base 12 and/or lid 14 can be fabricated of materials such as aluminum, stainless steel, and/or other materials. In exemplary embodiments lid 14 can be referred to as a body or  
35 as a flange and/or an adapter. According to exemplary aspects, base 12 can be configured to define at least a portion of volume 16. In exemplary embodiments walls 17

of base 12 can be affixed to lid 14 by a hermetical seal (not shown). An exemplary hermetical seal can include an O-ring arrangement between lid 14 and base 12. An appropriate removable fastener (not shown) may also be provided to maintain lid 14 and base 12 in a sealed arrangement.

5           In one embodiment, mass spectrometry assembly 10 comprises one or more components configured to perform operations with respect to mass spectrometry analysis, and accordingly, such components may be referred to as mass spectrometry components 30. In one possible implementation, lid 14 is coupled with one or more of components 30. Further, an individual one of components 30 may be internally or  
10           externally coupled with lid 14. For example, in the embodiment shown in Fig. 1, lid assembly 11 comprises lid 14 coupled with an external component 19 and an internal component 20. In the illustrated exemplary embodiment, external component 19 is external of a volume 16 and internal component 20 is at least partially within volume 16. An individual one of components 30 may also be provided coupled with lid 14 in other  
15           arrangements, for example, internal of or defined by a volume of lid 14.

          Within volume 16 at least some mass spectrometry operations can be performed using internal component 20. Some mass spectrometry operations can also be performed using external component 18. In an exemplary aspect, lid 14 and/or lid assembly 11 can be configured to be removably operably coupled with respect to base  
20           12. Lid 14 can be configured to be positioned in a first operable position 21. In position 21, lid 14 can form a hermetical seal with base 12 and provide component 20 within volume 16. In position 21, mass spectrometry assembly 10 can be used to perform at least some operations with respect to analysis of a sample.

          Lid 14 can also be configured to be positioned in a second operable position 23.  
25           In position 23, at least a portion of lid 14 can be spaced from base 12 and component 20 can be at least partially removed from volume 16. In an exemplary aspect, an entirety of lid 14 can be spaced from base 12 and component 20 can be at least partially removed from volume 16. In another exemplary aspect, at least a portion of lid 14 can be spaced from base 12 and component 20 can be entirely removed from volume 16. In another  
30           exemplary aspect, an entirety of lid 14 can be spaced from base 12 and an entirety of component 20 can be removed from volume 16. Second operable position 23 facilitates access to internal equipment 20 in one embodiment. Other operable positions intermediate operable positions 21 and 23 are possible. The plural operable positions of lid 14 and/or lid assembly 11 may refer to an exemplary embodiment of assembly 10  
35           where lid 14 and/or lid assembly 11 are detached and reattached numerous times with

respect to base 12 when used during mass spectrometry operations (e.g. service, reconfiguration, maintenance, etc.).

Referring next to Fig. 2, components 30 of a mass spectrometer according to one embodiment are shown. As represented in Fig. 2, components 30 can include a sample inlet component 32 operationally connected to an ion source component 34 which can be operationally connected to a mass separator component 35 which can be operationally connected to a detector component 36. These general components can be operationally connected to a processing and control device component 38. Exemplary embodiments provide for the use of components 30 to perform mass spectrometry. Components 30 can be operationally connected as shown in Fig. 2 or operationally connected in other configurations enabling mass spectrometry operations. Further, other arrangements including more or less or alternative components are possible.

As depicted in Fig. 2, a sample 40 can be introduced into sample inlet component 32. For purposes of this disclosure, sample 40 represents any chemical composition including both inorganic and organic substances in solid, liquid and/or vapor form. Specific examples of sample 40 suitable for analysis include volatile compounds such as toluene or other specific examples including highly-complex non-volatile protein based structures such as bradykinin. In certain aspects, sample 40 can be a mixture containing more than one substance or in other aspects sample 40 can be a substantially pure substance. Analysis of sample 40 can be performed according to exemplary aspects described below.

Sample inlet component 32 can be configured to introduce an amount of sample 40 into assembly 10 (Fig. 1) for analysis. Depending upon sample 40, sample inlet component 32 may be configured to prepare sample 40 for ionization. Types of sample inlet components 32 can include batch inlets, direct probe inlets, chromatographic inlets, and permeable, semi-permeable, solid phase microextraction (SPME), and/or capillary membrane inlets. Sample inlet component 32 can also include means for preparing sample 40 for analysis in the gas, liquid and/or solid phase. In some aspects, sample inlet component 32 may be combined with ion source component 34.

Ion source component 34 can be configured in exemplary embodiments to receive sample 40 directly or in other exemplary embodiments to receive sample 40 from sample inlet component 32. Ion source component 34 can be configured to convert portions or an entirety of sample 40 into analyte ions in one example. This conversion can include the bombardment of sample 40 with electrons, ions, molecules, and/or photons. This conversion can also be performed by thermal or electrical energy.

Ion source component 34 may utilize, for example, electron ionization (EI, typically suitable for the gas phase ionization), photo ionization (PI), chemical ionization, collisionally activated disassociation and/or electrospray ionization (ESI). For example in PI, the photo energy can be varied to vary the internal energy of the sample. Also, when  
5 utilizing ESI, sample 40 can be energized under atmospheric pressure and potentials applied when transporting ions into volume 16 of exemplary mass spectrometer assembly 10 (Fig. 1) can be varied to cause varying degrees of dissociation.

The analyte ions can proceed to mass separator component 35. Mass separator component 35 can include one or more of linear quadrupoles, triple quadrupoles,  
10 quadrupole ion traps (Paul), cylindrical ion traps, linear ion traps, rectilinear ion traps, ion cyclotron resonance, quadrupole ion trap/time-of-flight mass spectrometers, or other structures. Mass separator component 35 can also include focusing lenses as well as tandem mass separator components such as tandem ion traps or ion traps and quadrupoles in tandem. In one implementation at least one of multiple tandem mass  
15 separator components can be an ion trap. Tandem mass separator components can be placed in series or parallel. In an exemplary implementation, tandem mass separator components can receive ions from the same ion source component. In an exemplary aspect the tandem mass separator components may have the same or different geometric parameters. The tandem mass separator components may also receive  
20 analyte ions from the same or multiple ion source components.

Analytes may proceed to detector component 36. Exemplary detector components include electron multipliers, Faraday cup collectors, photographic and scintillation-type detectors. The progression of mass spectrometry analysis from sample inlet component 32 to detector component 36 can be controlled and monitored by a  
25 processing and control device component 38.

Acquisition and generation of data can be facilitated with processing and control device component 38. Processing and control device component 38 can be a computer or mini-computer or other appropriate circuitry that is capable of controlling components  
30 source component 34 and mass separator component 35, as well as the introduction of sample 40 via sample inlet component 32 and may further include determining, storing and ultimately displaying mass spectra recorded from detector component 36. Processing and control device component 38 can contain data acquisition and searching software. In one aspect such data acquisition and searching software can be configured  
35 to perform data acquisition and searching that includes the programmed acquisition of total analyte count. In another aspect, data acquisition and searching parameters can

include methods for correlating the amount of analytes generated to predetermine programs for acquiring data.

Referring again to Fig. 1, individual ones of the general components of a mass spectrometer may be positioned as an internal component 20 or as an external component 19 as desired by those of ordinary skill in the art. For example, in different applications or configurations, individual ones of general components 32-36 may be arranged as internal or external components as desired. In one embodiment, internal component(s) 20 can include one or more of sample inlet component 32, ion source component 34, mass separator component 35 and/or the detector component 36 in various configurations to perform mass spectrometry. In one embodiment, external component(s) 19 can include one or more of sample inlet component 32, ion source component 34 and/or processing and control device 36.

In an exemplary embodiment, internal and/or external components include multiple components such as multiple ion source components. These multiple components can be configured as external, internal or external and internal components.

Exemplary arrangements of the mass spectrometry components and lid assemblies are shown in Figs. 3a-3d.

Referring to Fig. 3a, an exemplary mass spectrometer assembly 10a is shown that includes lid 14a, base 12a and volume 16a. Lid 14a can include an interior surface 42a and an exterior surface 44a. Exemplary embodiments include lid assembly 11a that can include external component 19a including ion source component 34a coupled to external surface 44a. In the exemplary depiction, lid assembly 11a can include internal components 20a including sample inlet component 32a and mass separator component 35a including focusing lenses 50a and an ion trap 52a coupled to internal surface 42a to process analytes in a direction that is substantially perpendicular to the alignment of inner surface 42a toward the detector component (not shown). For example, analytes can be processed in a direction towards bottom 18a of base 12a. Sample inlet component 32a can be located between mass separator component 35a and internal surface 42a. Sample inlet component 34a and mass separator component 35a can be in a substantially stacked configuration below and coupled to inner surface 42a.

Referring to Fig. 3b, an exemplary mass spectrometry assembly 10b is shown that includes ion source component 34b coupled to external surface 44b. Lid assembly 11b can include lid 14b coupled to one or both of external component 19b and/or internal component 20b. As exemplarily depicted, internal components 20b include sample inlet component 34b and mass separator components 35b including focusing lenses 50b and ion trap 52b coupled to internal surface 42b to process analytes in a direction that is



substantially parallel to the alignment of inner surface 42b toward the detector component (not shown). For example, analytes can be processed in a direction towards wall 17b of base 12b.

Referring again to Figs. 3a and 3b, in operable position 21, internal components 20a-b including sample inlet component 34a-b, focusing lenses 50a-b and ion trap 52a-b and detector (not shown) can be within volume 16a-b. In an exemplary embodiment volume 16a-b can at least partially surround internal components 20a-b when lid 14a-b is in operable position 21. In other exemplary aspects internal components 20a-b can also include ion source component 34a-b.

In operable position 23, one or more of internal components 21a-b including sample inlet component 34a-b, focusing lenses 50a-b, and ion trap 52a-b can be at least partially removed from volume 16a-b. In an exemplary aspect, in operable position 23, one or more of the internal components can be entirely removed from volume 16a-b. For example and by way of example only, in operable position 23: sample inlet component 32a-b can be entirely removed from volume 16a-b while mass separator component 35a-b is not removed; sample inlet component 34a-b and focusing lenses 50a-b can be entirely removed from volume 16a-b while ion trap 52a-b is not removed; sample inlet component 32a-b and ion trap 52a-b can be entirely removed while focusing lenses 50a-b are not removed; both sample inlet component 32a-b and mass separator component 35a-b including both focusing lenses 50a-b and ion trap 52a-b can be entirely removed; focusing lenses 50a-b can be entirely removed while ion trap 52a-b is not removed; and/or ion trap 52a-b can be entirely removed while focusing lenses 50a-b are not removed from volume 16a-b.

Referring next to Fig. 3c, another exemplary mass spectrometry assembly 10c is shown and includes base 12c and lid 14c. As shown in Fig. 3c, lid 14c can include an interior surface 42c, an exterior surface 44c, and an opening 66 extending from interior surface 42c to exterior surface 44c. In the exemplary Figure, ion source component 34c is coupled to exterior surface 44c. In an exemplary aspect ion source component 34c can be coupled to exterior surface 44c in fluid communication with opening 66. In the exemplary depiction, internal components 20c can be coupled to interior surface 42c. In an exemplary aspect internal components 20c can couple to interior surface 42c in fluid communication with opening 66. Exemplary aspects include providing fluid communication between external components 19c and internal components 20c via opening 66. In an exemplary embodiment, ion source component 34c is in fluid communication with sample inlet component 32c through opening 66. Lid assembly 11c

can include lid 14c having opening 66 and one or both of internal components 20c and external components 19c.

Lid 14c can also be configured to provide sample 40 (Fig. 2) to sample inlet component 32c. In an exemplary embodiment openings 70 can be provided that extend  
5 into volume 16c from outside volume 16c through lid 14c. Sample tubing 72 can be provided through or as part of openings 70 and configured to provide sample 40 to sample inlet component 32c. In first operable position 21 internal component(s) 20c including sample inlet component 32c can be within volume 16c. In second operable position 23, portions or an entirety of one or more internal component(s) 20c which can  
10 include sample inlet component 32c can be removed from volume 16c.

Referring next to Fig. 3d, mass spectrometry assembly 10d is shown including lid 14d having an interior surface 42d and an exterior surface 44d having opening 66d therethrough, and edges 88 and 90 extending between interior surface 42d and exterior surface 44d. In an exemplary embodiment, lid 14d includes openings 92 and 94  
15 extending from edge 88 and edge 90 respectively to and in fluid communication with opening 66d. In an exemplary aspect openings 92 and 94 can provide fluid communication between outside of volume 16d and opening 66d. Conduit or tubing 96 can be configured to facilitate fluid communication to opening 66d through openings 92 and 94. Embodiments of lid 14d also include single openings extending from a single  
20 edge such as edge 88 and opening 92 to opening 66d. Embodiments of lid 14d also include openings (not shown) extending from outside volume 16d such as exterior surface 86 to opening 66. Openings 92 and 94 can be utilized in exemplary aspects to: provide auxiliary pumping to opening 66d; to provide reagent gas such as chemical ionization reagent gases to opening 66d during ionization using ion source 34d; and/or to  
25 provide sample 40 to opening 66d. In an exemplary embodiment, opening 92 can serve as an inlet port while opening 94 may serve as an outlet port for samples circulating through tubing within lid 14d. Openings 92 and/or 94 can also provide for gas pumping on central opening 66d should an intermediate pressure be needed for proper operation of ion source component 34d or in the event of a large sample influx such as is the case  
30 with liquid samples.

Lid 14d can also be configured to provide control and/or power to internal component 20d for example through electrical wiring 98. Electrical wiring 98 can be incorporated as part of lid 14d or through openings provided in lid 14d. Electrical wiring 98 can be configured to control internal component(s) 20d such as sample inlet  
35 component 32 and mass separator component 35 from processing and control device component 38. In first operable position 21 internal component(s) 20d and at least some

wiring 98 can be within volume 16d. In second operable position 23, portions or an entirety of one or more internal component(s) 20d and wiring 98 can be removed from volume 16d. Exemplary embodiments provide for lid assembly 11d that includes lid 14d and one or both of internal component 19d and/or internal component 20d

5 Referring next to Fig. 4, an exemplary mass spectrometer vacuum chamber lid assembly 11e is shown in isomeric view that includes a lid 14e and internal components 20e which may comprise mass separator components. Assembly 11e may be used in the arrangements of Figs. 1 and 3 in exemplary configurations. Components 20e can be coupled to lid 14e via one or more of a mounting rod 106 and a retainer clip 108. Mass  
10 separator components 20e can be confined within mass separator housings 110 and can be controlled via electrical signals provided to electrical connections 112. In one embodiment, mass separator housing 110 includes insulating material. As shown is Fig. 4, lid 11e includes openings 104 (see e.g., opening 70, Fig. 3c) and 102 (see e.g., opening 92, Fig 3d). In an exemplary aspect openings 104 can be used to pass wiring,  
15 sample, and/or other mass spectrometry components into volume 16 (Fig. 1). Opening 102 may be used as described above with reference to opening 92.

Referring next to Figs. 5a-5c an exemplary embodiment of lid 14f is shown in differing views. Referring first to Fig. 5a an exterior view of lid 14f is shown having recessed portions 114 radially inward of edges 116 providing an elevated lip 118.  
20 Recessed portion 114 can also be referred to as a groove. Openings 124 are provided through lid 14f in an exemplary embodiment to provide for feed through connections of electrical wiring and/or sample inlet components. Lid 14f can also include elevated portions 120, in an exemplary embodiment bisecting lid 14f. Opening 126 can be provided through the center of lid 14f to facilitate, in an exemplary aspect, the  
25 introduction of ions, samples or other materials provided during mass spectrometry. Lid 14f can also include adaptive portions 122 to facilitate the coupling of external components 19 (not shown) such as ion source component 34. In an exemplary embodiment opening 112 can extend bisecting lid 14f and through opening 126 to provide fluid communication between edge 116 and opening 126.

30 Referring next to Fig. 5b, internal surface view of body 14f is shown that includes a recessed portion 128 that can facilitate the hermetical sealing of lid 14f to base 12 of a mass spectrometry assembly 10 (Fig. 1). Referring next to Fig. 5c, a side view of lid 14f is shown. As shown lip 118 and adaptive portions 122 can extend above recessed surface 114.

35 Referring next to Fig. 6, an external view of an exemplary embodiment of lid 14g is shown that includes recessed groove 134 as well as a mating surface 132 to facilitate

sealing lid 14g to base 12 (Fig. 1). In an exemplary embodiment, base 12 can include a complimentary mating surface (not shown) to mating surface 132. Exemplary embodiments of lid 14g can also include a circular plate 136 that in certain aspects can be installed within opening 138 to maintain a pressure in opening 126 (Figs 5a-c) that is higher than that within volume 16 (Fig. 1).

Referring to Fig. 7, a mass spectrometry assembly 10f is shown that includes a base 142, a lid 144 and an external component 146. As shown, base portion 142 also includes an opening 148 that in an exemplary embodiment allows access to vacuum pumping and/or detection of ions separated utilizing assembly 10f. As exemplarily depicted in Fig. 7, external component 146 can include ion source component 34 and in one embodiment can be removably operably coupled with respect to lid 144 and configured to be positioned in a first operable position (not shown) to seal with lid 144 and a second operable position where at least a portion of external component 146 is spaced from lid 144.

Other aspects provide for the configuration of assembly 10f with multiple components. Multiple ion sources can be configured to couple with lid 144 in one embodiment. In an exemplary aspect, different ion sources can be configured to be exchanged and/or replaced with respect to assembly 10f. In an exemplary embodiment, an electron impact ion source may be replaced with a chemical ionization ion source.

Referring to the figures discussed above, mass spectrometer operational methods are also provided that include first performing mass analysis using mass spectrometry assembly 10 in operable position 21. This performance can include providing sample 40 to volume 16 as described above. According to an exemplary aspect, mass analysis can include providing ions to the vacuum chamber volume through opening 66 (Fig. 3c) and contacting the ions with sample 40. Contacting sample 40 with the ions can occur within opening 66. In an exemplary embodiment mass analysis can include providing a chemical ionization plasma and a chemical ionization reagent gas, for example through openings 92, to opening 66d (Fig. 3d) to produce ions and exposing sample 40 to the ions to form analytes.

After performing mass analysis, lid 14 can be moved to second operable position 23. In an exemplary aspect lid 14 can be at least partially removed from base 12 and internal component 20 can be at least partially removed from volume 16. During mass analysis, components 30 (Fig. 2) may require inspection between analysis for example for maintenance and trouble shooting requirements.

Internal components 20 such as mass separator components 35 (Fig. 2) can be inspected with lid 14 in second operable position 23. In an exemplary embodiment ion

source component 34, such as internal ion source components, including, in exemplary aspects, electron impact filaments, can become fouled during mass analysis and with lid 14 in second operable position 23, ion source components such as the filaments can be replaced with clean filaments or replacement filaments. In an exemplary embodiment  
5 mass separator component 35 such as ion trap 52a-b (Figs. 3a-b) can become fouled or require replacement during mass analysis and with lid 14 in second operable position 23 mass separator component 35 such as ion trap 52a-b can be replaced with a clean or conditioned ion trap. In an exemplary aspect sample inlet component 34, such as a semi-permeable membrane can require cleaning and or replacement during mass  
10 analysis and with lid 14 in second operable position 23 sample inlet component 32 can be cleaned or replaced.

In an exemplary embodiment, before moving the lid to operable position 23, ion source component 146 (Fig. 7) can be at least partially or entirely removed from lid 144 and inspected. Upon inspection, ion source component 146 can be cleaned, replaced,  
15 or otherwise manipulated.

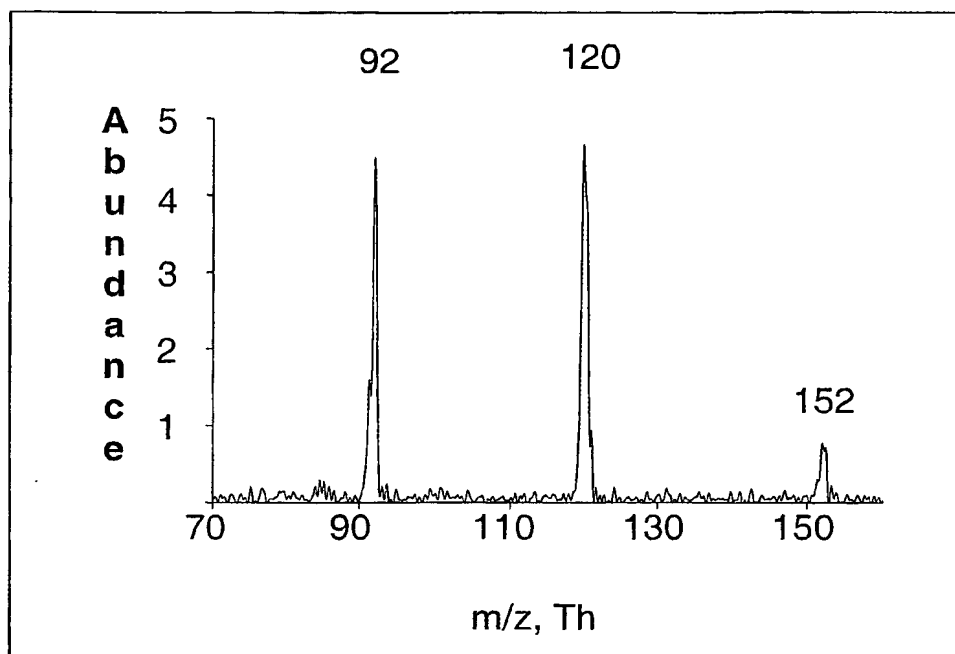
After inspection, the lid can be returned to first operable position (not shown) and mass analysis can be performed using components 30 (Fig. 2).

At least one arrangement facilitates servicing and reconfiguration of assembly 10. For example, upon removal of the lid assembly or the lid from the base, the internal  
20 components, wiring, and tubing, may be removed from the vacuum chamber thereby facilitating servicing, replacement, etc. of such components away from the confines of the vacuum chamber and perhaps reducing the chances of contamination. In one arrangement, the mere removal of the lid also removes at least one or more internal components in the same step. In other arrangements only internal components of  
25 interest are removed or perhaps even partially removed to facilitate inspection and/or maintenance while other internal components or portions of components of interest remain within the vacuum chamber. In one arrangement, the lid can be completely removed from the base of assembly 10 that may facilitate the inspection and maintenance of the internal components without the encumbrances of attachments to, or  
30 the confines of the base. In another arrangement, the external components can be removed from the lid to perhaps facilitate the inspection of the external component without substantially increasing the pressure within or contaminating the vacuum chamber. It is also contemplated that lid 14 or lid assembly 11 may remain partially coupled to base 12 in the second operable position (e.g. coupled via a hinge)

35 The following non-limiting examples are provided to further to facilitate aspects of the disclosure with respect to exemplary mass spectrometry operations of assembly 10.

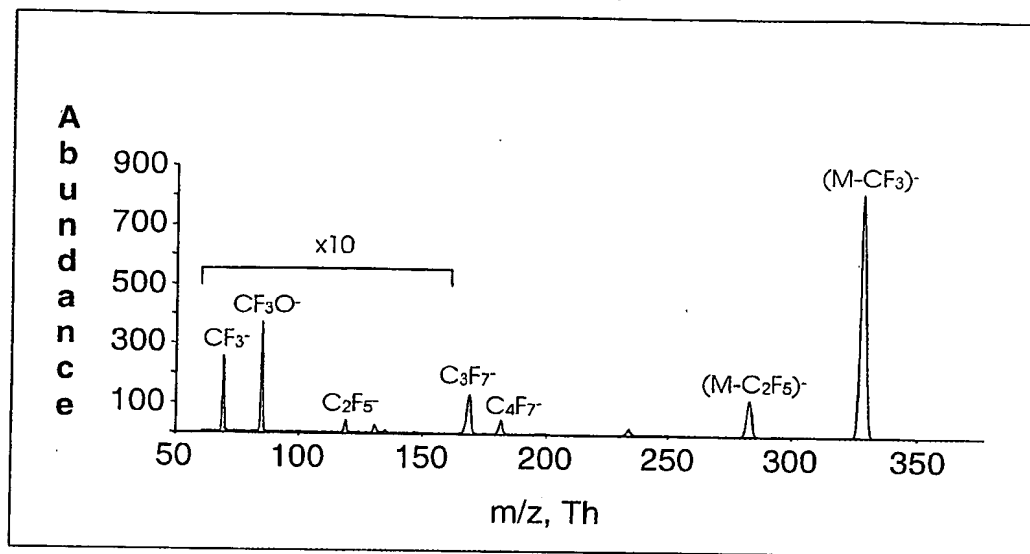
Methyl salicylate spectrum A below is obtained with internal Membrane Introduction Mass Spectrometry (MIMS)/internal Electron Ionization (EI) using a Griffin Analytical Technologies (West Lafayette, IN) Minotaur Model 2100A CIT Mass Spectrometer.

5

**Spectrum A**

Perfluorodimethylcyclohexane (PDCH) spectrum B below is obtained with direct inlet/external glow discharge ionization using a Griffin Analytical Technologies (West Lafayette, IN) Minotaur Model 2100A CIT Mass Spectrometer.

**Spectrum B**

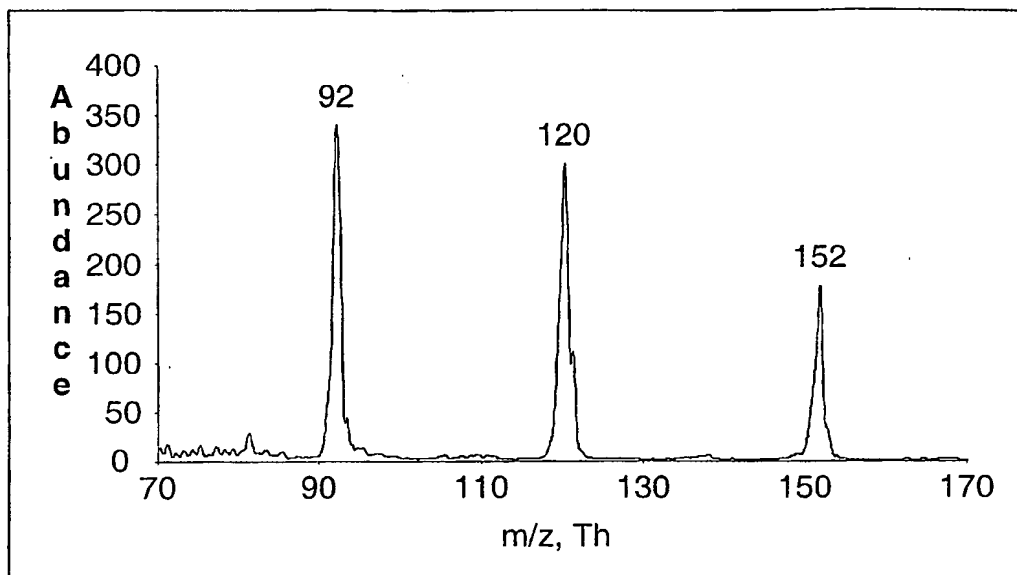


5

Note: spectrum was collected in two segments

Methyl salicylate spectrum C below is obtained with external Membrane Introduction Mass Spectrometry (MIMS)/internal Electron Ionization (EI) using a Griffin Analytical Technologies (West Lafayette, IN) Minotaur Model 2100A CIT Mass Spectrometer.

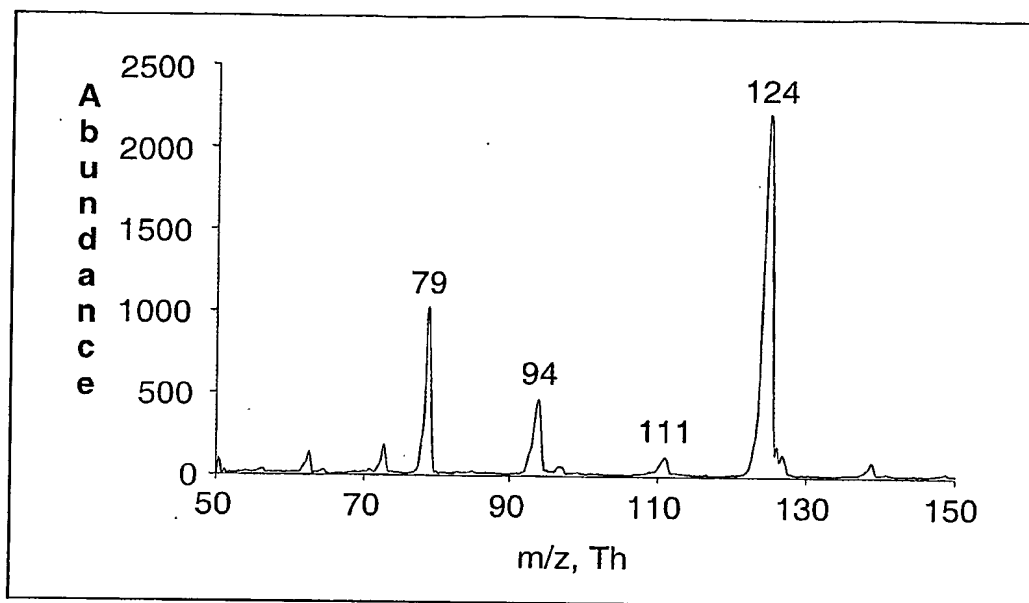
5

**Spectrum C**



Dimethyl Methylphosphonate (DMMP) spectrum D below is obtained with Solid-Phase Microextraction (SPME)/internal Electron Ionization (EI) using a Griffin Analytical Technologies (West Lafayette, IN) Minotaur Model 2100A CIT Mass Spectrometer.

**Spectrum D**



5

**CLAIMS**

What is claimed is:

1. A mass spectrometer assembly comprising:
  - 5 a base configured to define at least a portion of a vacuum chamber volume within which at least some operations may be performed with respect to mass spectrometry;
  - a mass separator component configured to perform at least some mass separation operations within the vacuum chamber volume; and
  - a lid coupled to the mass separator component and configured to be removably
  - 10 operably coupled with respect to the base, wherein the lid is configured to be positioned in a first operable position to form a hermetical seal with the base and provide the mass separator component at least partially within the vacuum chamber volume, and a second operable position wherein at least a portion of the lid is spaced from the base and the mass separator component is at least partially removed from the vacuum chamber
  - 15 volume.
2. The mass spectrometer assembly of claim 1 wherein the mass separator component comprises an entirety of the mass separator.
3. The mass spectrometer assembly of claim 1 wherein an entirety of the mass separator component is within the vacuum chamber volume.
- 20 4. The mass spectrometer assembly of claim 1 wherein the mass separator component is entirely removed from the vacuum chamber volume with the lid positioned in the second operating position.
5. The mass spectrometer assembly of claim 1 wherein an entirety of the lid is spaced from the base with the lid in the second operable position.
- 25 6. The mass spectrometer assembly of claim 1 wherein the mass separator component comprises at least one focusing lens.
7. The mass spectrometer assembly of claim 1 wherein the mass separator component comprises an ion trap.
8. The mass spectrometer assembly of claim 1 wherein the mass separator
- 30 component comprises focusing lenses and an ion trap.
9. The mass spectrometer assembly of claim 1 wherein the lid further comprises an opening configured to receive at least one electrical connection configured to connect to the mass separator component.

10. The mass spectrometer assembly of claim 1 wherein the lid further comprises an opening configured to introduce a sample to the vacuum chamber volume.
11. The mass spectrometer assembly of claim 1 wherein the lid further comprises an opening configured to introduce ions to the vacuum chamber volume.
- 5 12. The mass spectrometer assembly of claim 1 further comprising an external component coupled to the lid and configured to perform at least one operation with respect to mass spectrometry.
13. The mass spectrometer assembly of claim 12 wherein the external component comprises an ion source.
- 10 14. The mass spectrometer assembly of claim 12 wherein the external component comprises a plurality of ion sources.
15. The mass spectrometer assembly of claim 14 wherein one of the plurality of ion sources comprises an electron impact ion source and another of the plurality of ion sources comprises a chemical ionization ion source.
- 15 16. The mass spectrometer assembly of claim 12 wherein the external component comprises an inlet component.
17. The mass spectrometer assembly of claim 12 wherein the external component comprises both an inlet component and an ion source component.
- 20 18. A mass spectrometry vacuum chamber lid assembly comprising:  
a body;  
a mass separator component coupled to the body and configured to perform at least some operations with respect to mass separation for use during mass spectrometry; and  
wherein the body is configured to at least partially define a volume at least partially surrounding the mass separator component when the body is hermetically sealed to a base of a vacuum chamber assembly, wherein the body is further configured to be removable from the base to at least partially remove the mass separator component from the vacuum chamber volume.
- 25 19. The mass spectrometry vacuum chamber lid assembly of claim 18 wherein the mass separator component comprises at least one focusing lens.
- 30 20. The mass spectrometry vacuum chamber lid assembly of claim 18 wherein the mass separator component comprises an ion trap.

21. The mass spectrometry vacuum chamber lid assembly of claim 18 further comprising a sample inlet component coupled to the body.
22. The mass spectrometry vacuum chamber lid assembly of claim 21 wherein the sample inlet component comprises a semi-permeable membrane.
- 5 23. The mass spectrometry vacuum chamber lid assembly of claim 18 further comprising an ion source component coupled to the body.
24. The mass spectrometry vacuum chamber lid assembly of claim 23 wherein the body comprises an exterior surface and the ion source component is coupled to the exterior surface.
- 10 25. The mass spectrometry vacuum chamber lid assembly of claim 24 wherein the body further comprises an opening providing fluid communication between the ion source and the vacuum chamber volume and at least a portion of the sample inlet component is located between the opening and the mass separator component.
26. The mass spectrometry vacuum chamber lid assembly of claim 24
- 15 wherein the body further comprises a first opening providing fluid communication between the ion source component and the mass separator component.
27. The mass spectrometry vacuum chamber lid assembly of claim 26 wherein the body further comprises a second opening providing fluid communication between the first opening and outside the vacuum chamber volume.
- 20 28. The mass spectrometry vacuum chamber lid assembly of claim 27 wherein the body comprises at least one edge extending between exterior and interior surfaces and the second opening extends between the edge and the first opening, wherein the second opening is configured to provide one or more of reagent gas, sample, make up gas, and vacuum to the opening.
- 25 29. The mass spectrometry vacuum chamber lid assembly of claim 23 further comprising a sample inlet component coupled to an interior surface of the body.
30. The mass spectrometry vacuum chamber lid assembly of claim 29 wherein at least a portion of the sample inlet component is located between the interior surface and the mass separator component.

31. The mass spectrometry vacuum chamber lid assembly of claim 23 wherein the ion source component is configured to be removably operably coupled with respect to the lid and configured to be positioned in a first operable position to seal with the lid and a second operable position wherein at least a portion of the ion source component is spaced from the lid.

32. The mass spectrometry vacuum chamber lid assembly of claim 31 wherein the ion source is entirely removed from the lid in the second operable position.

33. The mass spectrometry vacuum chamber lid assembly of claim 18 wherein the mass separator component is configured to separate ions in a direction substantially parallel with the alignment of the interior surface of the body.

34. The mass spectrometry vacuum chamber lid assembly of claim 18 wherein the mass separator component is configured to separate ions in a direction substantially perpendicular with the alignment of the interior surface of the body.

35. The mass spectrometry vacuum chamber lid assembly of claim 18 further comprising an external component coupled to the body and configured to perform at least one operation with respect to mass spectrometry.

36. The mass spectrometry vacuum chamber lid assembly of claim 18 wherein the mass separator component comprises an entirety of the mass separator.

37. A mass spectrometer comprising:  
a vacuum chamber housing comprising a lid and a base, wherein the lid and the base are configured to define a vacuum chamber volume, wherein the base comprises a  
at least one wall configured to couple with the lid, wherein the lid comprises:  
5 an interior surface and an exterior surface;  
at least one edge extending between the interior and exterior surfaces;  
a first opening extending through the lid from the interior surface to the exterior surface; and  
a second opening extending from the edge to the first opening;  
10 a mass separator component coupled to the interior surface of the lid and configured to perform at least some operations with respect to mass separation for use in mass spectrometry;  
an ion source component coupled to the exterior surface of the lid and configured to perform at least some operations with respect to providing ions for use in mass  
15 spectrometry, wherein the first opening provides fluid communication between the mass separator and the ion source;  
wherein the lid is configured to be removably operably coupled with respect to the base and positioned in a first operable position to seal with the base and provide the mass separator component at least partially within the vacuum chamber volume, and a  
20 second operable position at least partially removed from the base to at least partially remove the mass separator component from the vacuum chamber volume; and  
a vacuum source in fluid communication with the vacuum chamber volume, wherein the seal of the base and the lid is configured to maintain a vacuum within the vacuum chamber volume sufficient to perform at least some operations with respect to  
25 mass spectrometry.
38. The mass spectrometry vacuum chamber lid assembly of claim 37 wherein an entirety of the mass separator component is with the vacuum chamber volume.
39. The mass spectrometry vacuum chamber lid assembly of claim 37  
30 wherein the seal comprises a hermetical seal.
40. The mass spectrometry vacuum chamber lid assembly of claim 37 wherein the mass separator component is entirely removed in the second operable position.

41. A mass spectrometer operational method comprising:  
providing a mass spectrometry assembly comprising a base, a lid and a mass separation component configured to perform at least some operations with respect to mass spectrometry, the base and lid substantially defining a vacuum chamber volume  
5 when the lid is affixed to the base, wherein the mass separator component is coupled to the lid and occupies a portion of the vacuum chamber volume with the lid affixed to the base;  
first performing mass analysis operations within the vacuum chamber volume using the mass separation component;  
10 after the first performing, at least partially removing the lid from the base, wherein the removing of the lid also at least partially removes the mass separator component from the vacuum chamber volume;  
inspecting the mass separator component with the mass separator component at least partially removed from the vacuum chamber volume;  
15 sealing the lid and the base after the inspecting; and  
second performing mass analysis operations using the mass separator component after the returning.
42. The mass spectrometer operational method of claim 41 wherein the removing entirely removes the lid from the base.
- 20 43. The mass spectrometer operational method of claim 41 wherein the removing entirely removes the mass separator component from the vacuum chamber volume.
44. The mass spectrometer operational method of claim 41 wherein the first performing mass analysis operations further comprises fouling the mass separator  
25 component and further comprising replacing the fouled mass separator component with a clean mass separator component.
45. The mass spectrometer operational method of claim 41 wherein the mass separator component comprises focusing lenses.
46. The mass spectrometer operational method of claim 41 wherein the mass  
30 separator component comprises an ion trap.
47. The mass spectrometer operational method of claim 41 wherein the mass separator component comprises focusing lenses and an ion trap.

48. The mass spectrometer operational method of claim 41 wherein the mass spectrometry assembly further comprises a sample inlet component coupled to the lid and the sample inlet component occupies a portion of the vacuum chamber volume with the lid and base affixed.

5 49. The mass spectrometer operational method of claim 48 wherein the sample inlet component comprises a semi-permeable membrane.

50. The mass spectrometer operational method of claim 41 wherein the mass spectrometry assembly further comprises an ion source component coupled to the lid and the ion source component occupies a portion of the vacuum chamber volume with  
10 the lid affixed to the base.

51. The mass spectrometer operational method of claim 50 wherein the ion source component comprises an electron impact ion source.

52. The mass spectrometer operational method of claim 41 wherein the mass spectrometry assembly further comprises an ion source component coupled to the lid  
15 and a sample inlet component coupled to the lid, and wherein the ion source and sample inlet components occupy a portion of the vacuum chamber volume with the lid affixed to the base.

53. The mass spectrometer operational method of claim 52 wherein the at least partially removing the lid also at least partially removes the ion source component  
20 and the sample inlet component from the vacuum chamber volume.

54. The mass spectrometer operational method of claim 41 wherein the mass spectrometry assembly further comprises an ion source component coupled to the lid and the ion source component occupies a portion of the vacuum chamber volume with the lid affixed to the base, and the first performing the mass analysis comprises providing  
25 ions from the ion source to the vacuum chamber volume.

55. The mass spectrometer operational method of claim 41 wherein the lid comprises an opening and the first performing the mass analysis comprises providing a chemical ionization plasma and a chemical ionization reagent gas to the vacuum chamber volume using the opening.

30 56. The mass spectrometer operational method of claim 41 wherein the first performing the mass analysis comprises providing ions to the vacuum chamber volume through an opening extending through the lid, providing sample to vacuum chamber volume, and contacting the ions with the sample.



57. The mass spectrometer operational method of claim 56 further comprising providing a first pressure within the opening and a second pressure within the vacuum chamber volume.

58. The mass spectrometer operational method of claim 57 wherein the first  
5 and second pressures are the same.

59. The mass spectrometer operational method of claim 57 wherein the first and second pressures are different.

60. The mass spectrometer operational method of claim 56 wherein the contacting the ions with sample occurs in the opening.

10 61. The mass spectrometer operational method of claim 41 wherein the mass spectrometry assembly further comprises an ion source component coupled to the exterior of the lid.

15 62. The mass spectrometer operational method of claim 61 wherein the ion source component comprises a plurality of ion sources and, before the at least partially removing the lid from the base, at least partially removing one of the plurality of ion sources from the lid.

63. The mass spectrometer operational method of claim 62 wherein, before the second performing, the one ion source is replaced with another ion source.

20 64. The mass spectrometer operational method of claim 61 wherein, before the at least partially removing the lid from the base, at least partially removing the ion source component from the lid.

65. The mass spectrometer operational method of claim 64 wherein, after at least partially removing the ion source component from the lid, inspecting the ion source component with the ion source component at least partially removed from the lid.

25 66. The mass spectrometer operational method of claim 61 wherein the first performing the mass analysis further comprises fouling the ion source component, and further comprising replacing the fouled ion source component with a clean ion source component.

30 67. The mass spectrometer operational method of claim 41 wherein the mass spectrometry assembly further comprises an external component.

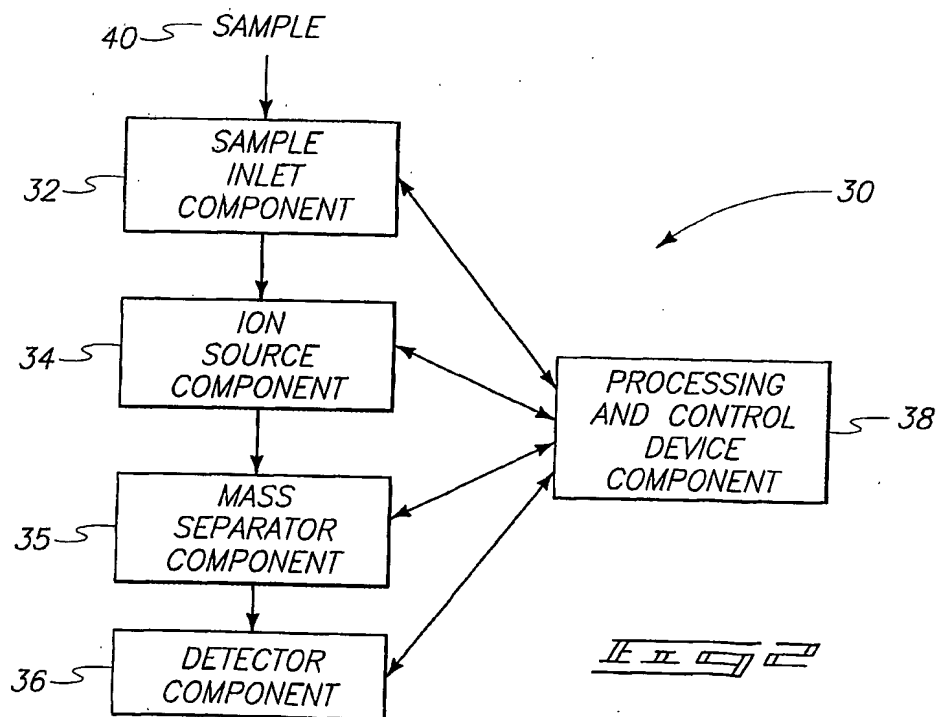
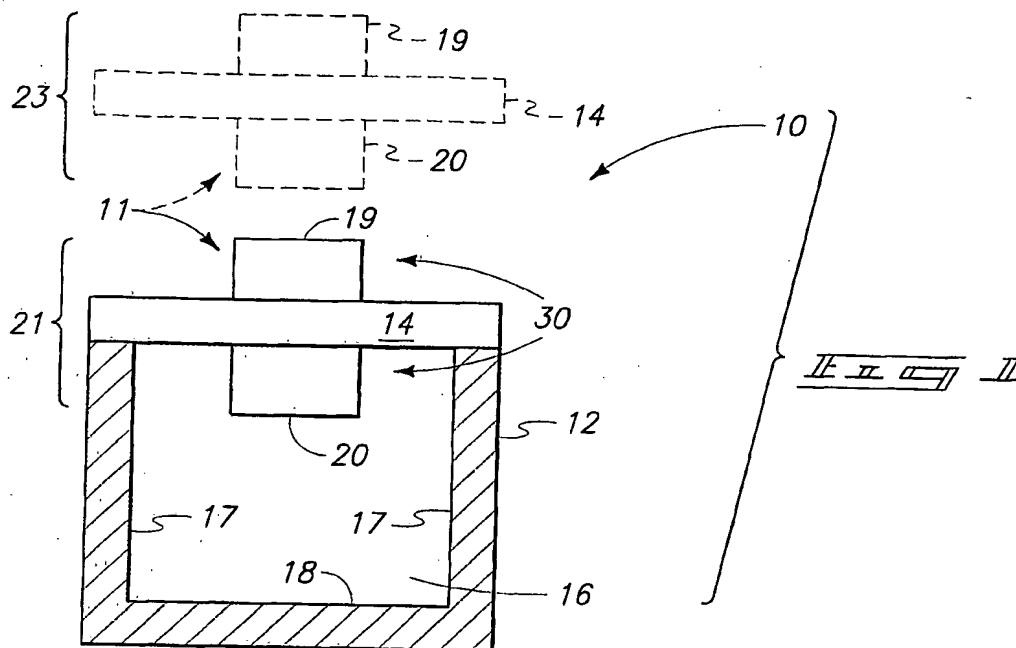
68. The mass spectrometer operational method of claim 67 wherein the external component comprises one or both of an ion source component and an sample inlet component.

5 69. The mass spectrometer operational method of claim 41 wherein the mass separator component comprises an entirety of the mass separator.

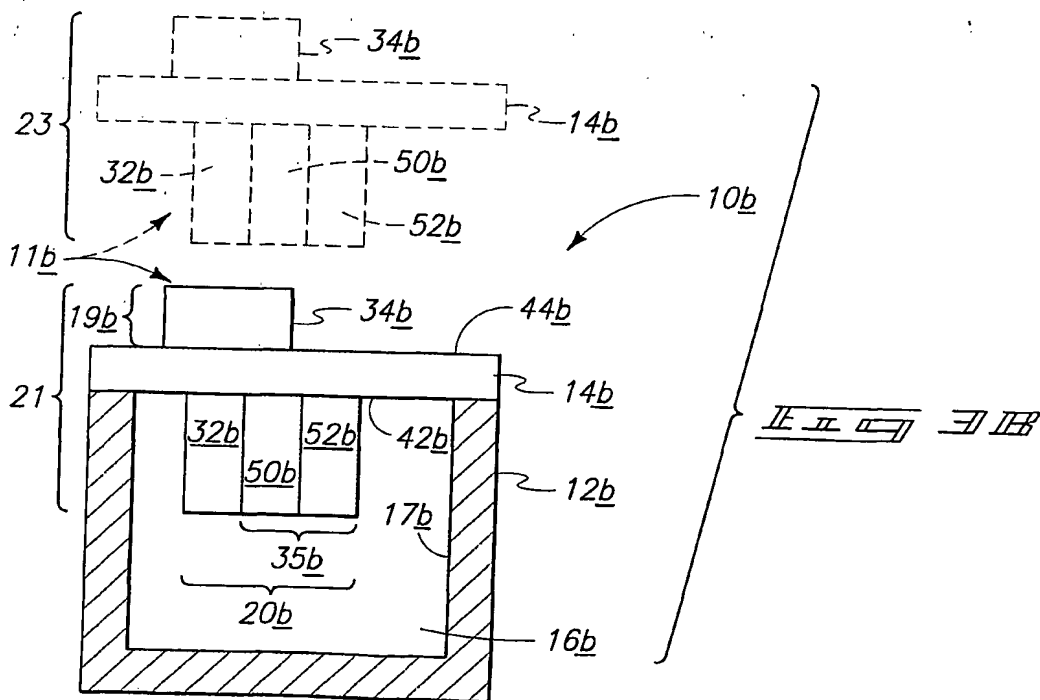
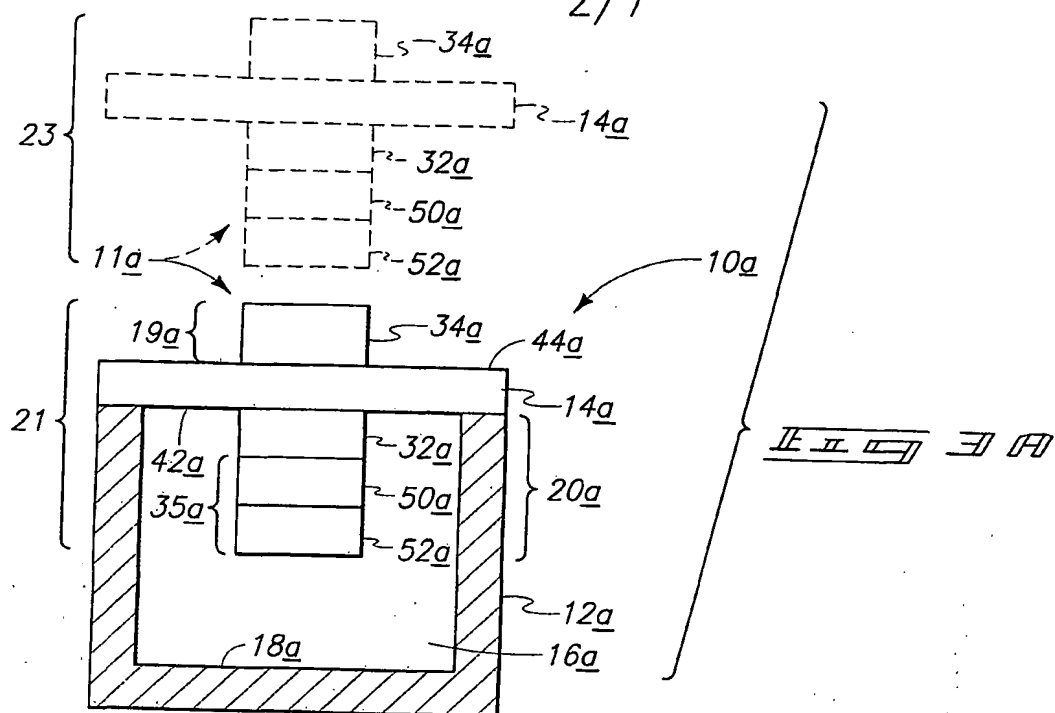
70. The mass spectrometer operational method of claim 67 wherein, before the second performing, exchanging the external component with another external component.

10 71. The mass spectrometer operational method of claim 70 wherein the external component comprises an electron impact ion source and the other external component comprises a chemical ionization ion source.

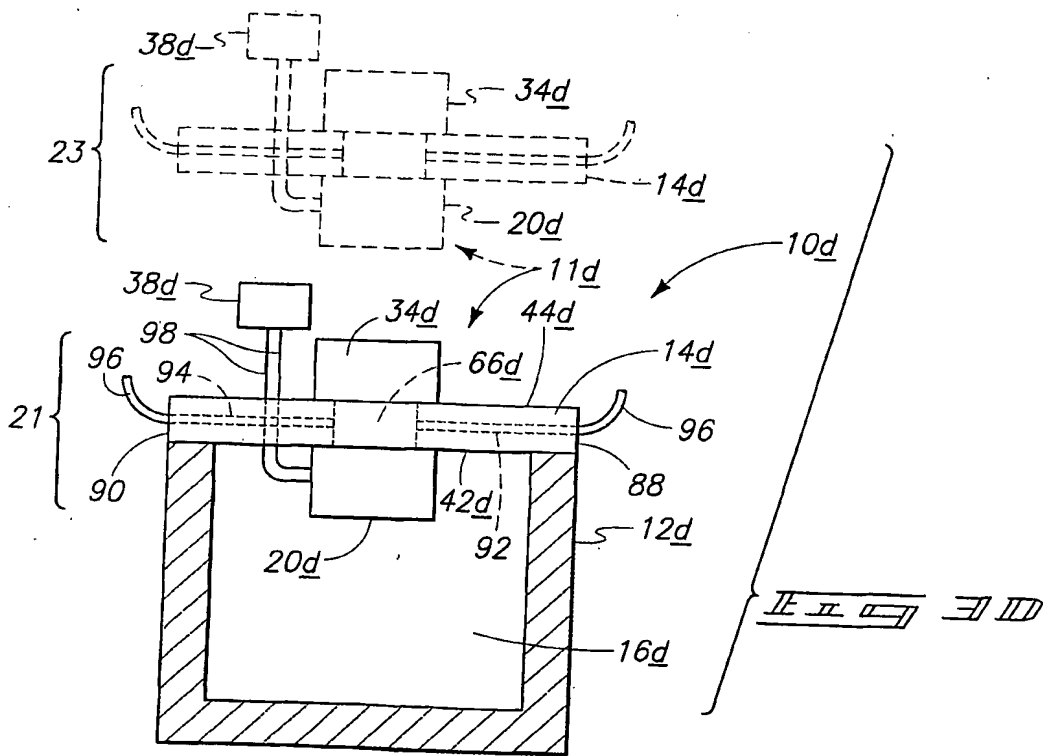
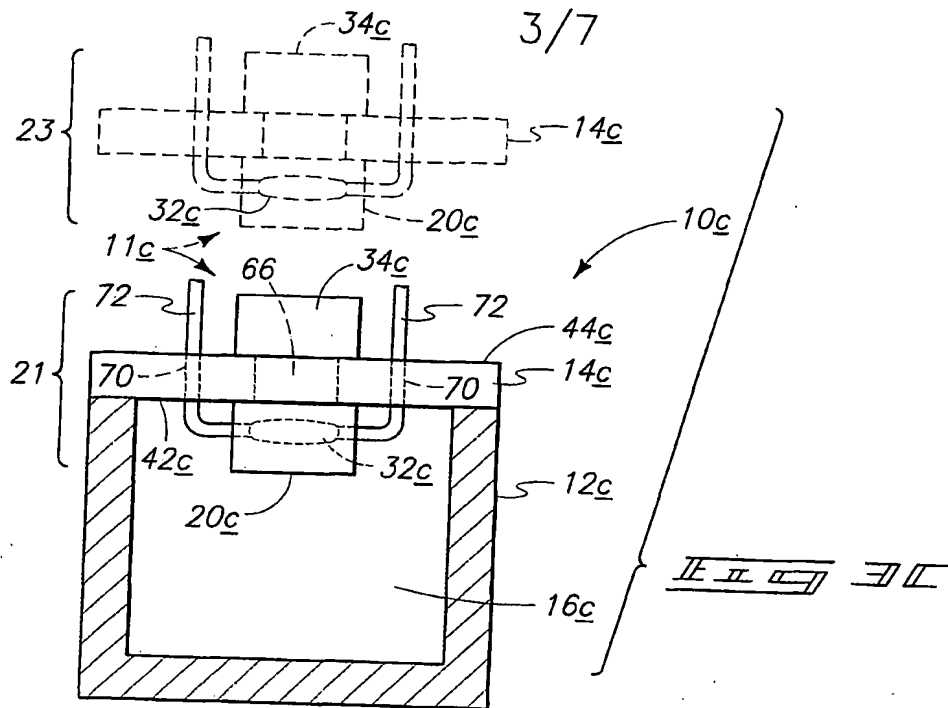
1/7



2/7



3/7



4/7

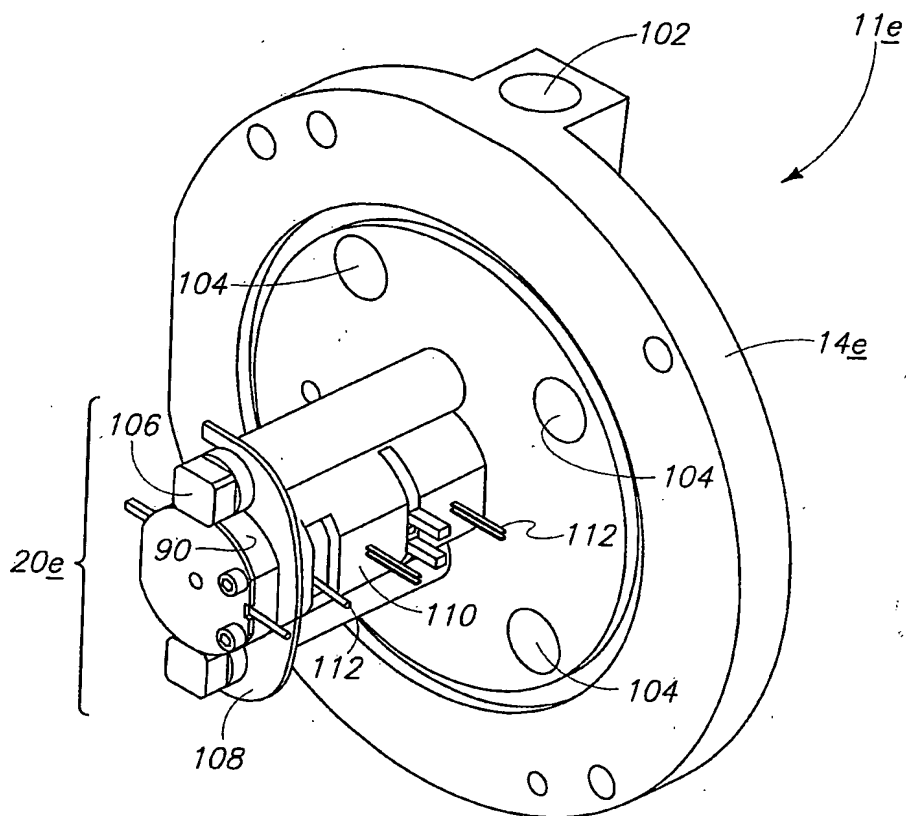
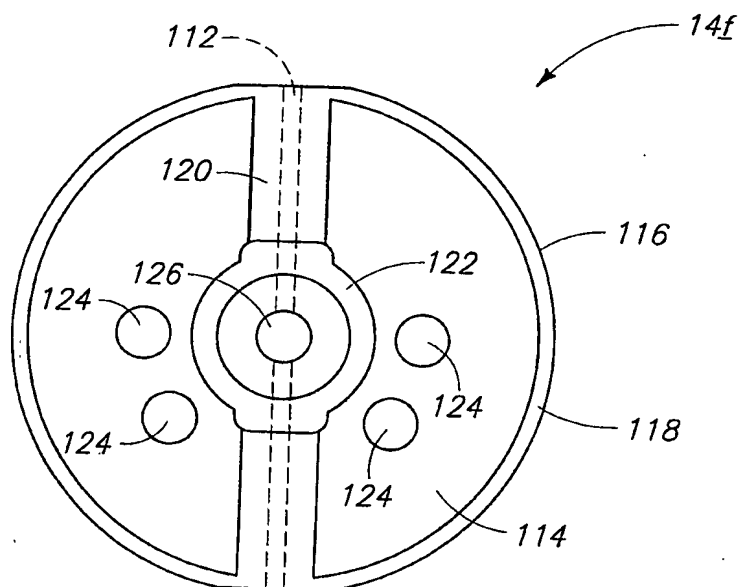
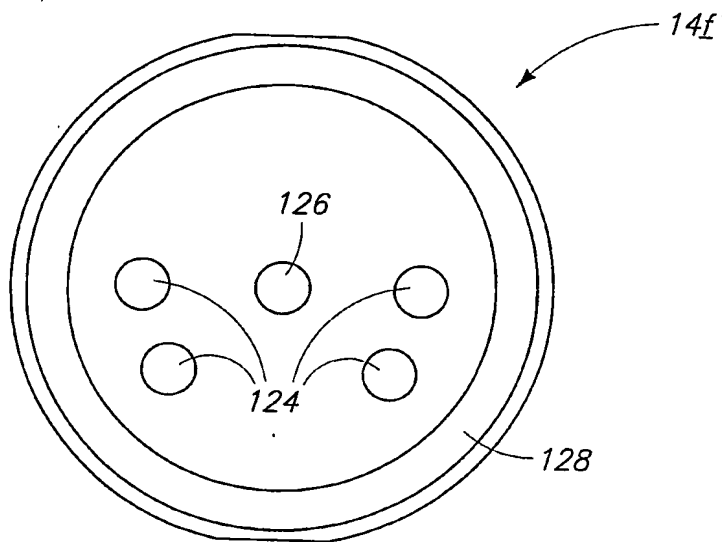


FIG. 4

5/7

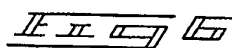
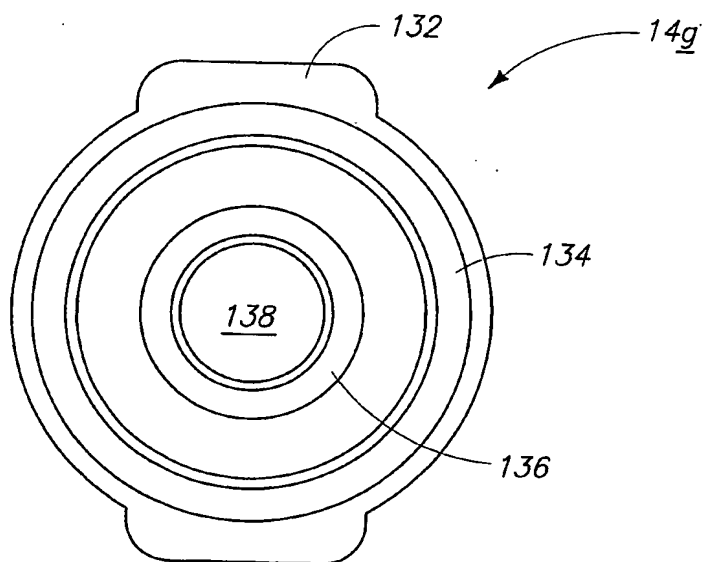
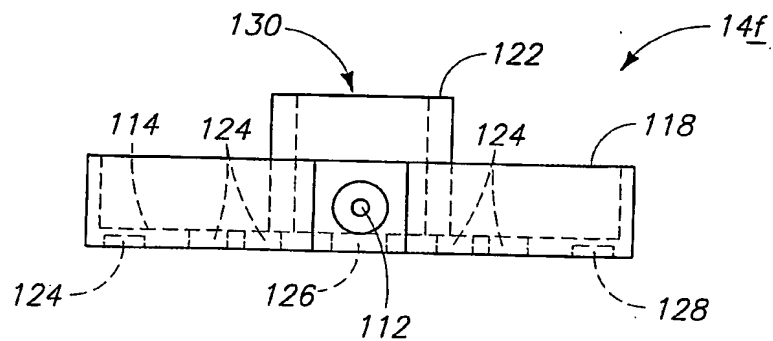


*FIG. 5A*



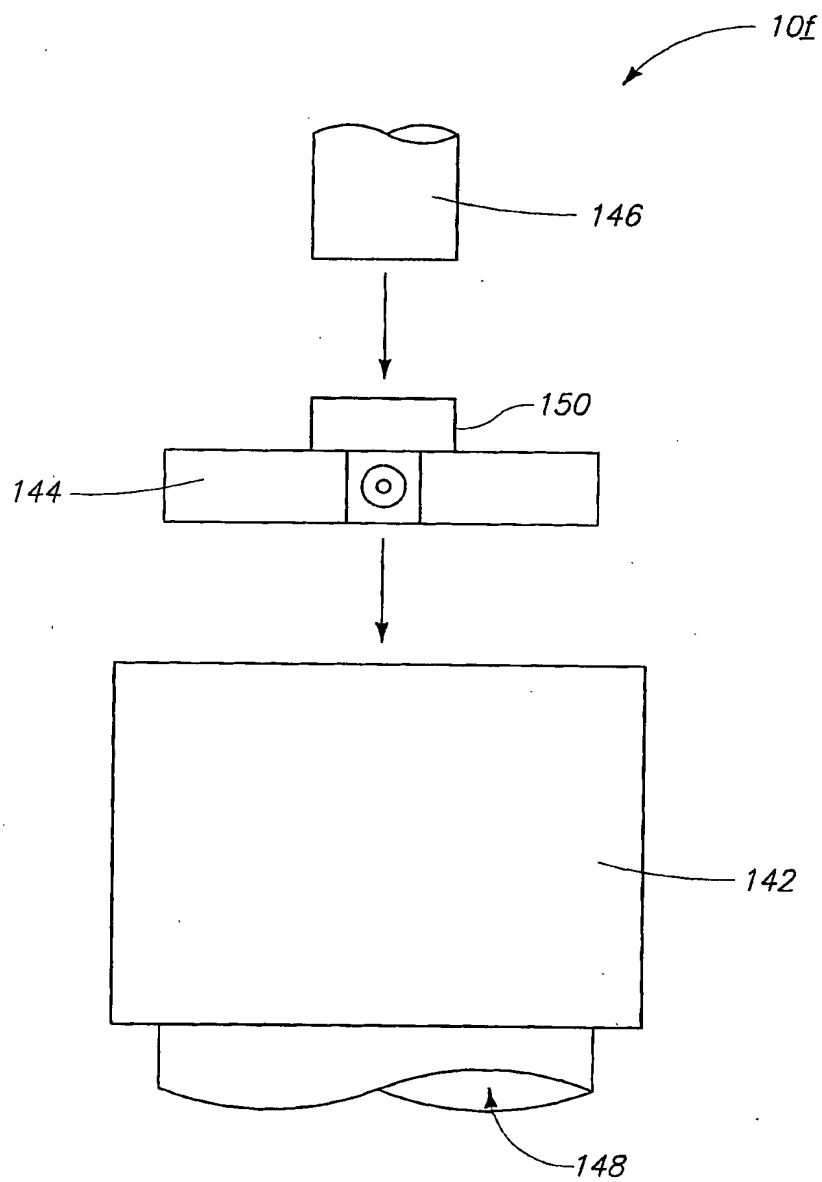
*FIG. 5B*

6/7





7/7



II II II II